## Amendments to the Claims

This listing of claims will replace all prior version and listings of claims in the application:

## Listing of Claims:

- (Currently amended): Combined pharmaceutical preparation for cancer therapy comprising as active [[substances]]substances:
  - a) at least one compound having glutaminase [[activity ]]activity; and
  - at least one antineoplastic agent selected from platinum complexes and anthracyclines.
- (Original): Preparation as claimed in claim 1, characterized in that the compound having glutaminase activity is a glutaminase, glutaminase-asparaginase, glutaminase analogue, derivative or modification of the same and is either of natural origin or is produced synthetically.
- (Original): Preparation as claimed in claim 2, characterized in that the compound having glutaminase activity is from Pseudomonas and is preferably Pseudomonas 7A glutaminase-asparaginase.
- (Currently amended): Preparation as claimed in <u>claim 1 one of the claims 1 to 3</u>, characterized in that the compound having glutaminase activity is modified preferably with polyethylene glycol.

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- (Currently amended): Preparation as claimed in <u>claim 1 one of the claims 1 to 4</u>, characterized in that it comprises doxorubicin, daunomycin, actinomycin D or/and mitoxantrone.
- (Currently amended): Preparation as claimed in <u>claim 1 one of the claims 1 to 5</u>, characterized in that
  it comprises cis-platinum, oxaliplatinum or/and carboplatinum.
- (Currently amended): Process for producing pharmaceutical preparations as claimed in <u>claim 1 one of the claims 1 to 6</u>, characterized in that the active substances optionally together with common pharmaceutical carrier substances or auxiliary substances are mixed and processed into oral or parenteral forms of administration.
- (Original): Use of in particular a compound having glutaminase activity and at least one antineoplastic agent selected from platinum complexes and anthracyclines to produce an agent for an antineoplastic therapy.
- abnormal cell proliferation,
  characterized in that
  at least one compound having glutaminase activity and at least one antineoplastic agent
  selected from platinum complexes or authracyclines are administered in a molar ratio

at least one compound naving gutaminase activity and at least one antineopiastic agent selected from platinum complexes or anthracyclines are administered in a molar ratio between 1:10 to 1:1000 and 10:1 to 1000:1, where the doses to be administered daily are 0.005 – 100 mg/kg body weight per individual component.

(Withdrawn): Method for treating cancer and other diseases which are associated with